limitations imposed upon such distribution by §1307.11(a)(4) and (b) of this chapter.

- (g) Owners or operators of vessels, aircraft, or other entities described in this section shall not be deemed to possess or dispense any controlled substance acquired, stored and dispensed in accordance with this section.
- (h) The Master of a vessel shall prepare a report for each calendar year which shall give in detail an accounting for all controlled substances purchased, dispensed, or disposed of during the year. The Master shall file this report with the medical officer emploxed by the owner or operator of his vessel, if any, or, if not, he shall maintain this report with other records required to be kept under the Act and, upon request, deliver a copy of the report to the Administration.
- (i) Controlled substances acquired and possessed in accordance with this section shall not be distributed to persons not under the general supervision of the medical officer employed by the owner or operator of the vessel, aircraft, or other entity, except in accordance with §1307.21 of this chapter.

[37 FR 15918, Aug. 8, 1972, as amended at 38 FR 756, Jan. 4, 1973. Redesignated at 38 FR 26609, Sept. 24, 1973, and amended at 41 FR 9546, Mar. 5, 1976; 50 FR 31589, Aug. 5, 1985]

## §1301.29 Provisional registration of narcotic treatment programs; compounders.

- (a) All persons currently approved by the Food and Drug Administration under §310.505 (formerly §130.44) of this title to conduct a methadone treatment program and who are registered by the Drug Enforcement Administration under this section will be granted a Provisional Narcotic Treatment Program Registration.
- (b) The provisions of §1301.45–1301.57 relating to revocation and suspension of registration, shall apply to a provisional registration.
- (c) Unless sooner revoked or suspended under paragraph (b) of this section, a provisional registration shall remain in effect until (1) the date on which such person has registered under this section or has had his registration denied, or (2) such date as may be prescribed by written notification to the

person from the Drug Enforcement Administration for the person to become registered to conduct a narcotic treatment program, whichever occurs first.

[39 FR 37984, Oct. 25, 1974]

## APPLICATIONS FOR REGISTRATION

## §1301.31 Time for application for registration; expiration date.

- (a) Any person who is required to be registered and who is not so registered may apply for registration at any time. No person required to be registered shall engage in any activity for which registration is required until the application for registration is granted and a Certificate of Registration is issued by the Administrator to such person.
- (b) Any person who is registered may apply to be reregistered not more than 60 days before the expiration date of his registration.
- (c) At the time a manufacturer, distributor, researcher, analytical lab, importer, exporter or narcotic treatment program is first registered, that business activity shall be assigned to one of twelve groups, which shall correspond to the months of the year. The expiration date of the registrations of all registrants within any group will be the last date of the month designated for that group. In assigning any of the above business activities to a group, the Administration may select a group the expiration date of which is less than one year from the date such business activity was registered. If the business activity is assigned to a group which has an expiration date less than three months from the date of which the business activity is registered, the registration shall not expire until one year from that expiration date; in all other cases, the registration shall expire on the expiration date following the date on which the business activity is registered.
- (d) At the time a retail pharmacy, hospital/clinic, practitioner or teaching institution is first registered, that business activity shall be assigned to one of twelve groups, which shall correspond to the months of the year. The expiration date of the registrations of all registrants within any group will be the last day of the month designated for that group. In assigning any of the

above business activities to a group, the Administration may select a group the expiration date of which is not less than 28 months nor more than 39 months from the date such business activity was registered. After the initial registration period, the registration shall expire 36 months from the initial expiration date.

[36 FR 7778, Apr. 24, 1971, as amended at 36 FR 13386, July 21, 1971. Redesignated at 38 FR 26609, Sept. 24, 1973, and amended at 52 FR 20599, June 2, 1987]

## §1301.32 Application forms; contents; signature.

- (a) If any person is required to be registered, and is not so registered and is applying for registration:
- (1) To manufacture or distribute controlled substances, he shall apply on DEA Form 225:
- (2) To dispense controlled substances listed in Schedules II through V, he shall apply on DEA Form 224;
- (3) To conduct instructional activities with controlled substances listed in Schedules II through V, he shall apply on DEA Form 224;
- (4) To conduct research with controlled substances listed in Schedules II through V (other than research described in §§1301.22(a)(6), he shall apply on DEA Form 225;
- (5) To conduct research with narcotic drugs listed in Schedules II through V, as described in §1301.22(a)(6), he shall apply on DEA Form 225;
- (6) To conduct research with controlled substances listed in Schedule I, he shall apply on DEA Form 225, with three copies of a research protocol as described in § 301.33(a) attached to the form, or, in the case of a clinical investigation, with three copies of a certificate of submission of an IND as described in §1301.33(b) attached to the form (the researcher also submitting to the Food and Drug Administration three copies of a Notice of Claimed Investigational Exemption for a New Drug as required in §1301.33(b));
- (7) To conduct instructional activities with controlled substances listed in Schedule I, he shall apply as a researcher on DEA Form 225 with two copies of a statement describing the nature, extent, and duration of such in-

structional activities attached to the form:

- (8) To conduct chemical analysis with controlled substances listed in any Schedule, he shall apply on DEA Form 225; and
- (9) To conduct a narcotic treatment program, including a compounder, shall apply on DEA Form 363.
- (b) If any person is registered and is applying for reregistration:
- (1) To manufacture or distribute controlled substances, he shall apply on DEA Form 225a;
- (2) To dispense controlled substances listed in Schedules II through V, he shall apply on DEA Form 224a;
- (3) To conduct instructional activities with controlled substances listed i Schedules II through V, he shall apply on DEA Form 224a;
- (4) To conduct research with controlled substances listed in Schedules II through V (other than research described in §1301.22(a) (6), he shall apply on DEA Form 225a;
- (5) To conduct research with narcotic drugs listed in Schedules II through V, as described in §1301.22(a) (6), he shall apply on DEA Form 225a;
- (6) To continue to conduct research with controlled substances listed in Schedule I under one or more approved research protocols, he shall apply on DEA Form 225a;
- (7) To continue to conduct instructional activities with controlled substances listed in Schedule I under one or more approved instructional statements, he shall apply as a researcher on DEA Form 225a;
- (8) To conduct chemical analysis with controlled substances listed in any Schedule, he shall apply on DEA Form 225a; and
- (9) To conduct a narcotic treatment program, including a compounder, shall apply on DEA Form 363a (Renewal Form).
- (c) DEA (or BND) Forms 224 and 225 may be obtained at any regional office of the Administration or by writing to the Registration Unit, Drug Enforcement Administration, Department of Justice, Post Office Box 28083, Central Station, Washington, DC 20005. DEA Forms 224a, 225a and 363a will be mailed, as applicable, to each registered person approximately 60 days